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(54) Title: NEW CRYSTALLINE FORM OF OMEPRAZOLE			
(57) Abstract  The present invention relates to a novel crystalline form of 5-methoxy-2-[[[4-methoxy-3,5-dimethyl- 2-pyridinyl)methyl] sulfinyl]-1H-benzimidazole, know under the generic name omeprazole. Further, the present invention also relates to the use of the novel crystalline form of 5-methoxy-2-[[[4-methoxy-3,5-dimethyl- 2-pyridinyl)methyl] sulfinyl]-1H-benzimidazole for the treatment of gastrointestinal disorders, pharmaceutical compositions containing it as well as processes for the preparation of the novel crystalline form of 5-methoxy-2-[[[4-methoxy-3,5-dimethyl- 2-pyridinyl)methyl] sulfinyl]-1H-benzimidazole.			

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## NEW CRYSTALLINE FORM OF OMEPRAZOLE

*Field of the invention*

5 The present invention relates to a novel crystalline form of 5-methoxy-2-[[[4-methoxy-3,5-dimethyl-2-pyridinyl)methyl]sulfinyl]-1H-benzimidazole. 5-methoxy-2-[[[4-methoxy-3,5-dimethyl-2-pyridinyl)methyl]sulfinyl]-1H-benzimidazole is known under the generic name omeprazole and its novel crystalline form is hereinafter referred to as omeprazole form A. Further, the present invention also relates to use of omeprazole form A for the treatment of  
10 gastrointestinal disorders, pharmaceutical compositions containing omeprazole form A and processes for the preparation of omeprazole form A.

*Background of the invention and prior art*

15 The compound 5-methoxy-2-[[[4-methoxy-3,5-dimethyl-2-pyridinyl)methyl]sulfinyl]-1H-benzimidazole, having the generic name omeprazole, as well as therapeutically acceptable salts thereof, are described in EP 5129. The single crystal X-ray data and the derived molecular structure of the so far only known crystal form of omeprazole is described by Ohishi *et al.*, Acta Cryst. (1989), C45, 1921-1923. This published crystal form of  
20 omeprazole is hereinafter referred to as omeprazole form B.

Omeprazole is a proton pump inhibitor, *i.e.* effective in inhibiting gastric acid secretion, and is useful as an antiulcer agent. In a more general sense, omeprazole may be used for treatment of gastric-acid related diseases in mammals and especially in man.

25

*Brief description of the drawings*

Figure 1 is an X-ray powder diffractogram of omeprazole form A.

Figure 2 is an X-ray powder diffractogram of omeprazole form B.

*Description of the invention*

It has surprisingly been found that the substance omeprazole can exist in more than one  
5 crystal form. It is an object of the present invention to provide omeprazole form A. Another  
object of the present invention is to provide a process for the preparation of omeprazole  
form A, substantially free from other forms of omeprazole. X-ray powder diffraction  
(XRPD) is used as a method of differentiating omeprazole form A from other crystalline  
and non-crystalline forms of omeprazole. Additionally it is an object of the present  
10 invention to provide pharmaceutical formulations comprising omeprazole form A.

Omeprazole form A is a crystalline form exhibiting advantageous properties, such as being  
well-defined, being thermodynamically more stable and less hygroscopic than omeprazole  
form B, especially at room temperature. Omeprazole form A does also show a better  
15 chemical stability, such as thermo stability and light stability, than omeprazole form B.

Omeprazole form B can under certain conditions, completely or partly, be converted into  
omeprazole form A. Omeprazole form A is thereby characterized in being thermo-  
dynamically more stable than omeprazole form B.

20

Omeprazole form A is further characterized as being essentially non-hygroscopic.

Omeprazole form A is characterized by the positions and intensities of the peaks in the X-  
ray powder diffractogram, as well as by the unit cell parameters. The unit cell dimensions  
25 have been calculated from accurate Guinier data. The X-ray powder diffractogram data as  
well as the unit cell parameters for omeprazole form B are different compared to  
omeprazole form A. Omeprazole form A can thereby be distinguished from omeprazole  
form B, using X-ray powder diffraction.

Omeprazole form A, according to the present invention, is characterized in providing an X-ray powder diffraction pattern, as in figure 1, exhibiting substantially the following d-values and intensities;

Form A		Form A	
d-value (Å)	Relative intensity	d-value (Å)	Relative intensity
9.5	vs	3.71	s
7.9	s	3.59	m
7.4	w	3.48	m
7.2	vs	3.45	s
6.0	m	3.31	w
5.6	s	3.22	s
5.2	s	3.17	m
5.1	s	3.11	w
4.89	w	3.04	w
4.64	m	3.00	w
4.60	m	2.91	w
4.53	w	2.86	w
4.49	m	2.85	w
4.31	m	2.75	w
4.19	w	2.67	w
4.15	w	2.45	w
3.95	w	2.41	w

The peaks, identified with d-values calculated from the Bragg formula and intensities, have been extracted from the Guinier diffractogram of omeprazole form A. The relative

intensities are less reliable and instead of numerical values the following definitions are used;

	% Relative Intensity*	Definition
5	25-100	vs (very strong)
	10-25	s (strong)
	3-10	m (medium)
	1-3	w (weak)

\* The relative intensities are derived from diffractograms measured with fixed slits.

10

Omeprazole form A according to the present invention is further characterized by a triclinic unit cell with parameters;

$$a=10.410(4) \text{ \AA}$$

$$b=10.468(3) \text{ \AA}$$

$$c=9.729(4) \text{ \AA}$$

$$\alpha=111.51(3)^\circ$$

$$\beta=116.78(3)^\circ$$

$$\gamma=90.77(3)^\circ$$

15 Omeprazole form A can also be characterized by Raman spectroscopy, where omeprazole form A is characterized by the absence of a band at  $1364 \text{ cm}^{-1}$ , which is observed for omeprazole form B, and by the ratio of the relative intensities of the  $842 \text{ cm}^{-1}$  and  $836 \text{ cm}^{-1}$  bands. The ratio (intensity of  $842 \text{ cm}^{-1}$  band / intensity of  $836 \text{ cm}^{-1}$  band) is  $<1$  for omeprazole form A, while the ratio is  $>1$  for omeprazole form B.

20

According to the invention there is further provided a process for the preparation of omeprazole form A.

Omeprazole form A is obtained upon slow crystallization and omeprazole form B is obtained from fast crystallization. Omeprazole form A may be prepared by reaction crystallisation or recrystallizing omeprazole of any form, or mixtures of any forms, in an appropriate solvent, such as for instance methanol, at around room temperature and for a  
5 prolonged time period. Examples of prolonged time periods include, but are not limited to, a few hours, such as 2 hours, up to several weeks. Suitable solvents are alkyl alcohols and especially a lower alcohol comprising 1 - 4 carbon atoms.

Omeprazole form A may also be prepared by suspending omeprazole of any form, or  
10 mixtures of any forms, in an appropriate solvent at around room temperature and for a prolonged time period. Examples of appropriate solvents include, but are not limited to, methanol, ethanol, acetone, ethyl acetate, methyl tert. butyl ether, toluene, or any mixture thereof. Examples of prolonged time periods include, but are not limited to, a few hours, such as 2 hours, up to several weeks.

15

The omeprazole form A obtained according to the present invention is substantially free from other crystal and non-crystal forms of omeprazole, such as omeprazole form B. Substantially free from other forms of omeprazole shall be understood to mean that omeprazole form A contains less than 10%, preferably less than 5%, of any other forms of  
20 omeprazole, *e.g.* omeprazole form B.

Omeprazole form A in mixture with other solid form/forms of omeprazole, *e.g.* omeprazole form B, also exhibits advantageous properties, such as being chemically more stable than pure omeprazole form B. Mixtures comprising a certain amount of omeprazole form A, by  
25 weight, are also chemically more stable than other mixtures comprising a lesser amount of omeprazole form A, by weight. Such mixtures comprising omeprazole form A can be prepared, for example, by mixing omeprazole form A prepared according to the present invention with other solid forms of omeprazole, such as form B, prepared according to prior art.

30

The present invention also relates to mixtures comprising omeprazole form A in mixture with other solid forms of omeprazole. Such mixtures comprising omeprazole form A include for instance mixtures containing a detectable amount of omeprazole form A, 1%, 2%, 5%, 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, 98% or 99% (by weight), of omeprazole form A.

Examples of other solid forms of omeprazole include, but are not limited to, omeprazole form B, amorphous forms, and other polymorphs.

A detectable amount of omeprazole form A is an amount that can be detected using conventional techniques, such as FT-IR, Raman spectroscopy, XRPD and the like.

The expression chemical stability includes, but is not limited to, thermo stability and light stability.

The compound of the invention, *i.e.* omeprazole form A, prepared according to the present invention is analyzed, characterized and differentiated from omeprazole form B by X-ray powder diffraction, a technique which is known per se. Another suitable technique to analyze, characterize and differentiate omeprazole form A from omeprazole form B is by Raman spectroscopy.

Omeprazole form A is effective as a gastric acid secretion inhibitor, and is useful as an antiulcer agent. In a more general sense, it can be used for treatment of gastric-acid related conditions in mammals and especially in man, including *e.g.* reflux esophagitis, gastritis, duodenitis, gastric ulcer and duodenal ulcer. Furthermore, it may be used for treatment of other gastrointestinal disorders where gastric acid inhibitory effect is desirable *e.g.* in patients on NSAID therapy, in patients with Non Ulcer Dyspepsia, in patients with symptomatic gastro-esophageal reflux disease, and in patients with gastrinomas. The compound of the invention may also be used in patients in intensive care situations, in patients with acute upper gastrointestinal bleeding, pre- and postoperatively to prevent



aspiration of gastric acid and to treat stress ulceration. Further, the compound of the invention may be useful in the treatment of psoriasis as well as in the treatment of *Helicobacter* infections and diseases related to these. The compound of the invention may also be used for treatment of inflammatory conditions in mammals, including man.

5

Any suitable route of administration may be employed for providing the patient with an effective dosage of omeprazole form A according to the invention. For example, peroral or parenteral formulations and the like may be employed. Dosage forms include capsules, tablets, dispersions, suspensions and the like, *e.g.* enteric-coated capsules and/or tablets, capsules and/or tablets containing enteric-coated pellets of omeprazole. In all dosage forms omeprazole form A can be admixed with other suitable constituents. .

10

According to the invention there is further provided a pharmaceutical composition comprising omeprazole form A, as active ingredient, in association with a pharmaceutically acceptable carrier, diluent or excipient and optionally other therapeutic ingredients. Compositions comprising other therapeutic ingredients are especially of interest in the treatment of *Helicobacter* infections. The invention also provides the use of omeprazole form A in the manufacture of a medicament for use in the treatment of a gastric-acid related condition and a method of treating a gastric-acid related condition which method comprises administering to a subject suffering from said condition a therapeutically effective amount of omeprazole form A.

15

20

The compositions of the invention include compositions suitable for peroral or parenteral administration. The compositions may be conveniently presented in unit dosage forms, and prepared by any methods known in the art of pharmacy.

25

In the practice of the invention, the most suitable route of administration as well as the magnitude of a therapeutic dose of omeprazole form A in any given case will depend on the nature and severity of the disease to be treated. The dose, and dose frequency, may also vary according to the age, body weight, and response of the individual patient. Special

30

requirements may be needed for patients having Zollinger-Ellison syndrome, such as a need for higher doses than the average patient. Children and patients with liver diseases as well as patients under long term treatment will generally benefit from doses that are somewhat lower than the average. Thus, in some conditions it may be necessary to use  
5 doses outside the ranges stated below. Such higher and lower doses are within the scope of the present invention.

In general, a suitable oral dosage form may cover a dose range from 5 mg to 250 mg total daily dose, administered in one single dose or equally divided doses. A preferred dosage  
10 range is from 10 mg to 80 mg.

The compound of the invention may be combined as the active component in intimate admixture with a pharmaceutical carrier according to conventional techniques, such as the oral formulations described in WO 96/01623 and EP 247 983, the disclosures of which are  
15 hereby incorporated as a whole by reference.

Combination therapies comprising omeprazole form A and other active ingredients in separate dosage forms, or in one fixed dosage form, may also be used. Examples of such active ingredients include anti-bacterial compounds, non-steroidal anti-inflammatory  
20 agents, antacid agents, alginates and prokinetic agents.

The examples which follow will further illustrate the preparation of the compound of the invention, i.e. omeprazole form A, but are not intended to limit the scope of the invention as defined hereinabove or as claimed below.

25

### *Examples*

#### Example 1

*Preparation of omeprazole form A*

Omeprazole (55.8 g) is added at room temperature to methanol (348 ml) containing ammonia (1.3 ml; 25%). The suspension is thereafter stirred in darkness for approximately  
5 45 hours and then filtered. The filtrate is dried 18 hours at 30°C under reduced pressure (<5 mbar). Yield: 43.9 g.

Example 210 *Preparation of omeprazole form B*

Omeprazole (50 g) is added to methanol (750 ml) containing ammonia (0.7 ml; 25%) at 50°C. The solution is thereafter filtered and cooled in about 20 minutes to approximately 0°C. The formed crystals are filtered and washed with ice cooled methanol and then dried.  
15 The filtrate was dried 24 hours at 40°C under reduced pressure (<5 mbar). Yield: 39 g.

Example 320 *Characterization of omeprazole form A and omeprazole form B using X-ray powder diffraction*

X-ray diffraction analysis was performed according to standard methods which can be found in e.g. Bunn, C. W. (1948), Chemical Crystallography, Clarendon Press, London; or Klug, H.P. & Alexander, L. E. (1974), X-Ray Diffraction Procedures, John Wiley and  
25 Sons, New York. The unit cell parameters for omeprazole form A and B have been calculated from the Guinier X-ray powder diffractograms using the program "TREOR" by Werner, P.-E., Eriksson, L. and Westdahl, M., J. Appl. Crystallogr. 18 (1985) 367 - 370. The fact that the positions of all peaks in the diffractograms for omeprazole form A and form B may be calculated using the respective unit cell parameters, proves that the unit cells are

correct and that the diffractograms are indicative of the pure forms. The diffractogram of omeprazole form A, prepared according to Example 1 in the present application, is shown in Figure 1 and the diffractogram of omeprazole form B, prepared according to Example 2 in the present application is shown in Figure 2.

5

The peaks, identified with d-values calculated from the Bragg formula and intensities, have been extracted from the diffractograms for omeprazole forms A and form B, and are given in Table 1. In this table the unit cell parameters for omeprazole forms A and B are also given. The relative intensities are less reliable and instead of numerical values the

10 following definitions are used;

% Relative Intensity	Definition
25-100	vs (very strong)
10-25	s (strong)
15 3-10	m (medium)
1-3	w (weak)

Some additional weak or very weak peaks found in the diffractograms have been omitted from table 1.

20

Table 1. X-ray powder diffraction data for omeprazole form A and form B shown in Figures 1 and 2. All peaks noted for omeprazole form A and form B can be indexed with the unit cells given below.

Form A		Form B	
d-value (Å)	Relative intensity	d-value (Å)	Relative intensity
9.5	vs	9.6	vs
7.9	s	8.0	m
7.4	w	7.9	m
7.2	vs	7.5	w
6.0	m	7.1	vs
5.6	s	5.9	m
5.2	s	5.6	m
5.1	s	5.3	s
4.89	w	5.1	s
4.64	m	4.54	m
4.60	m	4.48	s
4.53	w	4.41	m
4.49	m	4.14	w
4.31	m	3.75	s
4.19	w	3.57	m
4.15	w	3.47	s
3.95	w	3.40	w
3.71	s	3.28	s
3.59	m	3.22	m
3.48	m	3.02	w
3.45	s	2.97	w

Form A		Form B	
d-value (Å)	Relative intensity	d-value (Å)	Relative intensity
3.31	w	2.87	w
3.22	s	2.37	w
3.17	m		
3.11	w		
3.04	w		
3.00	w		
2.91	w		
2.86	w		
2.85	w		
2.75	w		
2.67	w		
2.45	w		
2.41	w		

The triclinic unit cells are:

Unit cell form A

$a=10.410(4) \text{ Å}$

$b=10.468(3) \text{ Å}$

$c=9.729(4) \text{ Å}$

$\alpha=111.51(3)^\circ$

$\beta=116.78(3)^\circ$

$\gamma=90.77(3)^\circ$

Unit cell form B

$a=10.257(10) \text{ Å}$

$b=10.717(6) \text{ Å}$

$c=9.694(10) \text{ Å}$

$\alpha=112.14(7)^\circ$

$\beta=115.56(5)^\circ$

$\gamma=91.76(7)^\circ$

**CLAIMS**

1. Omeprazole form A, characterized in being thermodynamically stable at room temperature.
2. Omeprazole form A, characterized in being essentially non-hygroscopic.
3. Omeprazole form A according to claims 1 or 2, characterized in providing an X-ray powder diffraction pattern exhibiting substantially the following d-values;

<b>Form A</b>		<b>Form A</b>	
d-value (Å)	Relative intensity	d-value (Å)	Relative intensity
9.5	vs	3.71	s
7.9	s	3.59	m
7.4	w	3.48	m
7.2	vs	3.45	s
6.0	m	3.31	w
5.6	s	3.22	s
5.2	s	3.17	m
5.1	s	3.11	w
4.89	w	3.04	w
4.64	m	3.00	w
4.60	m	2.91	w
4.53	w	2.86	w
4.49	m	2.85	w
4.31	m	2.75	w
4.19	w	2.67	w
4.15	w	2.45	w
3.95	w	2.41	w

4. Omeprazole form A, according to any of claims 1-3, characterized by having a triclinic unit cell with parameters
- 5       $a=10.410(4) \text{ \AA}$ ,  $b=10.468(3) \text{ \AA}$ ,  $c=9.729(4) \text{ \AA}$ ,  $\alpha=111.51(3)^\circ$ ,  $\beta=116.78(3)^\circ$ ,  
 $\gamma=90.77(3)^\circ$ .
5. Omeprazole, characterized in containing more than 50%, by weight, of omeprazole form A according to any of claims 1-4.
- 10
6. A process for the preparation of omeprazole form A as defined in any of claims 1-4, comprising the steps of;
- a) dissolving or suspending omeprazole of any form, or a mixture of any form, in a suitable solvent;
- 15      b) allowing the solution to crystallize, optionally using omeprazole form A to induce crystallization, and
- c) isolating the omeprazole form A thus obtained.
7. A process according to claim 6, characterized in that the solvent used in step a) is
- 20      chosen from a group consisting of methanol, ethanol, acetone, ethyl acetate, methyl tert. butyl ether, toluene, or any mixture thereof.
8. A process according to claims 6 or 7, characterized in that step a) is performed at 15-25°C.
- 25
9. A process according to any of claims 6-8, characterized in that step b) is performed during a prolonged time period.



10. A process according to any claim 6-9, characterized in that step b) is performed during at least 2 hours.

11. Omeprazole form A, prepared by a process according to any of claims 6-10.

5

12. A pharmaceutical formulation comprising omeprazole as defined in any of claims 1-5 in admixture with a pharmaceutically acceptable excipient.

13. The use of omeprazole as defined in any of claims 1-5, as active ingredient in the manufacture of medicament for use in treatment of gastrointestinal disorders.

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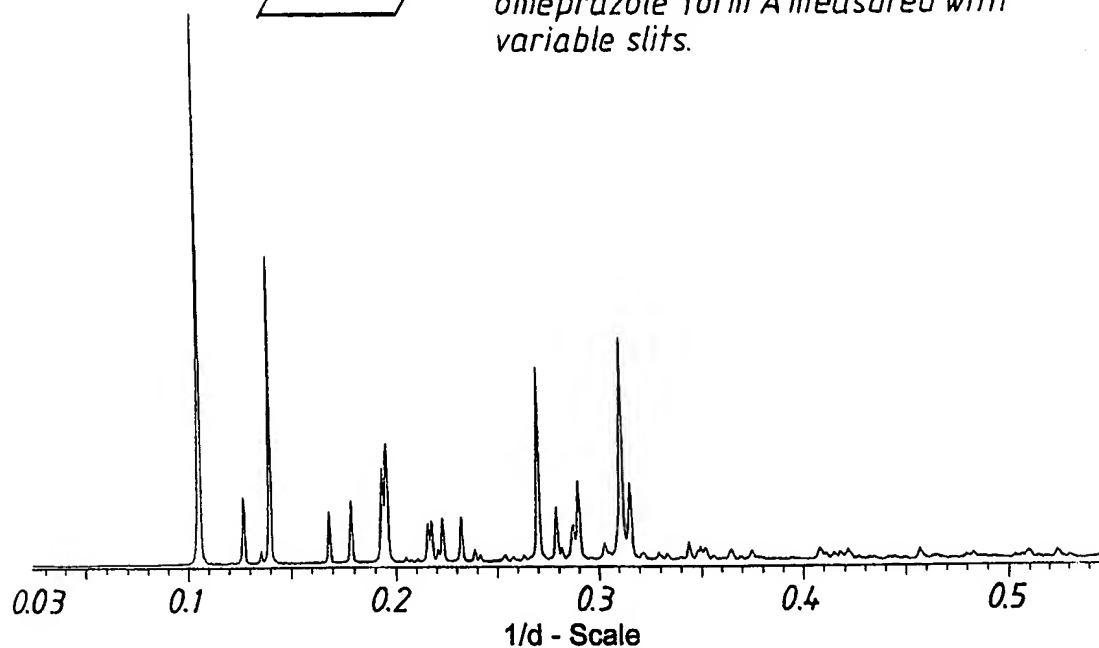
14. A method of treatment of gastrointestinal disorders which comprises administration of a therapeutically effective amount of omeprazole as defined in any of claims 1-5, to a patient suffering from gastrointestinal disorders.

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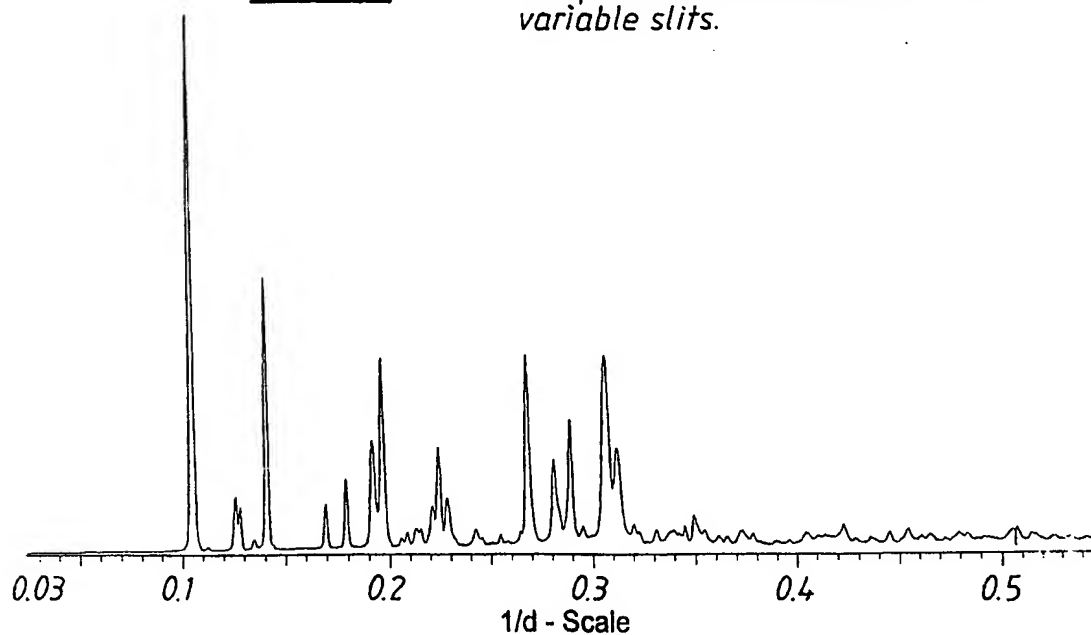
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**Fig. 1**

An X-ray powder diffractogram of omeprazole form A measured with variable slits.

**Fig. 2**

An X-ray powder diffractogram of omeprazole form B measured with variable slits.



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